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STATES OF NEW		Washington, D.C. 20231		

ATTORNEY DOCKET NO. APPLICATION NO. FILING DATE FIRST NAMED INVENTOR 89/520,091 CWIRLA 9 0300-0014 EXAMINER HM22/1002

REED & ASSOICIATES 3282 ALPHINE ROAD FORTOLA VALLEY CA 94028

SISSON, 3 ART UNIT PAPER NUMBER

DATE MAILED:

10/02/01

Please find below and/or attached an Office communication concerning this application or proceeding.

Commissioner of Patents and Trademarks

		Application N		Applicant(s)	
•			о.	'' ''	
0	Office Action Summary	09/620,091		CWIRLA ET AL.	
, O		Examiner		Art Unit	
The	MAILING DATE of this communica	Bradley L Siss		1655	ldroce
Period for Rep		nuon appears on the co	ver sneet with the t	correspondence ad	uress
THE MAILIN - Extensions of after SIX (6) M - If the period for Fallure to repl - Any reply rece	NED STATUTORY PERIOD FOR NO DATE OF THIS COMMUNITY in many be available under the provisions of 6 CONTROL from the partial but and the provisions of 6 CONTROL from the mailing date of this command or reply is appendix above, the maximum statut within the set or extended period for reply will wise by the Office later than three months after term adjustment. See 37 GFR 1,704(b).	ATION. 87 CFR 1.136(a). In no event, h cation. lays, a reply within the statutory bry period will apply and will exp , by statute, cause the applicatic	owever, may a reply be tin minimum of thirty (30) day ire SIX (6) MONTHS from in to become ABANDONE	nely filed s will be considered timel the mailing date of this o D (35 U.S.C. § 133).	y. ommunication.
1) Resp	oonsive to communication(s) filed	on			
2a) This	action is FINAL. 2b)☐ This action is nor	-final.		
	e this application is in condition fo ed in accordance with the practice				e merits is
Disposition of	Claims				
4)⊠ Claim	(s) 1-161 is/are pending in the ap	pplication.			
4a) Of	the above claim(s) is/are	withdrawn from consid	eration.		
5) Claim	(s) is/are allowed.				
6)☐ Claim	(s) is/are rejected.				
7)☐ Claim	(s) is/are objected to.				
8)⊠ Claim	(s) <u>1-161</u> are subject to restriction	n and/or election requi	rement.		
Application Pa	pers				
9)☐ The sp	ecification is objected to by the E	xaminer.			
10) The dr	awing(s) filed on is/are: a)	accepted or b) obje	ected to by the Exa	miner.	
Appli	cant may not request that any object	ion to the drawing(s) be l	neld in abeyance. S	ee 37 CFR 1.85(a).	
11) The pr	oposed drawing correction filed o	n is: a) 🗌 appro	ved b) disappro	ved by the Examin	er.
If app	proved, corrected drawings are requir	red in reply to this Office	action.		
12) ☐ The oa	th or declaration is objected to by	the Examiner.			
Priority under	35 U.S.C. §§ 119 and 120				
13) Ackno	wledgment is made of a claim for	r foreign priority under	35 U.S.C. § 119(a)-(d) or (f).	
a)∏ All	b)☐ Some * c)☐ None of:				
1.	Certified copies of the priority do	cuments have been re	ceived.		
2.	2. Certified copies of the priority documents have been received in Application No				
	Copies of the certified copies of tapplication from the Internation	onal Bureau (PCT Rul	∋ 17.2(a)).		Stage
	attached detailed Office action for				application)
a) 🗌 Th	rledgment is made of a claim for one translation of the foreign langu	age provisional applica	ation has been rec	eived.	аррисацоп).
	rledgment is made of a claim for	domestic priority unde	- 35 U.S.C. §§ 120	and/or 121.	
Attachment(s)	07. 1.000	г	7		
2) Notice of Draf	erences Cited (PTO-892) ftsperson's Patent Drawing Review (PTO- isclosure Statement(s) (PTO-1449) Papel		Interview Summary Notice of Informal F Other: Notice to Co		

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Election/Restrictions

- 1. Restriction to one of the following inventions is required under 35 U.S.C. 121:
 - Claims 1-16, 23-35, 42-53, 60-67, 74-93, 100-117, 124-140, and 147-155, drawn to a peptide compound, classified in class 530, subclass 328; and claims 17, 36, 54, 68, 94, 118, 141, and 156, drawn to a pharmaceutical composition comprising said peptide, classified in class 514, subclass 2.
 - II. Claims 18-22, 37-41, 55-59, 69-73, 95-99, 119-123, 142-146, and 157-161, drawn to a method of treating, classified in class 514, subclass 2.

The inventions are distinct, each from the other because of the following reasons:

- 2. Inventions I and II are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP § 806.05(h)). In the instant case the peptides as claimed can be used in an immunoassay.
- Because these inventions are distinct for the reasons given above and have acquired a separate status in the art as shown by their different classification, restriction for examination purposes as indicated is proper.

Sequence Restriction Requirement Applicable to Groups I and II

4. In addition the inventions detailed above read on patentably distinct Groups drawn to multiple nucleic acid fragments and/or polypeptide fragments found in multiple SEQ ID Numbers. The sequences are patentably distinct because they are unrelated sequences, and a Application/Control Number: 09/620,091

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further restriction is applied to each Group. For an elected Group drawn peptides or methods of using any of the polypeptide fragments, Applicants are permitted to elect <u>a single sequence</u> (See MPEP 803.04).

- Applicant is advised that the reply to this requirement to be complete must include an
 election of the invention to be examined even though the requirement be traversed (37 CFR
 1.143).
- 6. Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a petition under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Sequence Rules Compliance

7. This application contains sequence disclosures that are encompassed by the definitions for nucleotide and/or amino acid sequences set forth in 37 CFR 1.821(a)(1) and (a)(2). However, this application fails to comply with the requirements of 37 CFR 1.821 through 1.825 for the reason(s) set forth on the attached Notice To Comply With Requirements For Patent Applications Containing Nucleotide Sequence and/or Amino Acid Sequence Disclosures. Applicant must comply with the requirements of the sequence rules (37 CFR 1.821 - 1.825) before the application can be examined under 35 U.S.C. §8 131 and 132.

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8. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Bradley L Sisson whose telephone number is (703) 308-3978. The examiner can normally be reached on 6:30 a.m. to 5 p.m., Monday through Thursday.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, W. Gary Jones can be reached on (703) 308-1152. The fax phone numbers for the organization where this application or proceeding is assigned are (703) 305-3592 for regular communications and (703) 308-0294 for After Final communications.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (703) 308-0196.

Bradley L Sisson Primary Examiner Art Unit 1655

B. L. Sim

bls October 1, 2001

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NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

Applicant must file the items indicated below within the time period set the Office action to which the Notice is attached to avoid abandonment under 35 U.S.C. § 133 (extensions of time may be obtained under the provisions of 37 CFR 1.136(a)).

The nucleotide and/or amino acid sequence disclosure contained in this application does not comply with the requirements for such a disclosure as set forth in 37 C.F.R. 1.821 - 1.825 for the following reason(s):

1. This application clearly fails to comply with the requirements of 37 C.F.R. 1.821-1.825. Applicant's attention is directed to the final rulemaking notice published at 55 FR 18230 (May 1, 1990), if the effective filing date is on or after July 1, 1998, see the final rulemaking notice published at 63 FR 29620 (June 1, 1998) and 1211 OG 82 (June 23, 1998).
2. This application does not contain, as a separate part of the disclosure on paper copy, a "Sequence Listing" as required by 37 C.F.R. 1.821(c).
3, A copy of the "Sequence Listing" in computer readable form has not been submitted as required by 37 C.F.R. 1.821(e).
4. A copy of the "Sequence Listing" in computer readable form has been submitted. However, the content of the computer readable form does not comply with the requirements of 37 C.F.R. 1.822 and/or 1.823, as indicated on the attached copy of the marked -up "Raw Sequence Listing."
5. The computer readable form that has been filed with this application has been found to be damaged and/or unreadable as indicated on the attached CRF Diskette Problem Report. A Substitute computer readable form must be submitted as required by 37 C.F.R. 1.825(d).
6. The paper copy of the "Sequence Listing" is not the same as the computer readable from of the "Sequence Listing" as required by 37 C.F.R. 1.821(e).
7. Other:
Applicant Must Provide:
An initial or <u>substitute</u> computer readable form (CRF) copy of the "Sequence Listing".
An initial or <u>substitute</u> paper copy of the "Sequence Listing", as well as an amendment directing its entry into the specification.
A statement that the content of the paper and computer readable copies are the same and, where applicable, include no new matter, as required by 37 C.F.R. 1.821(e) or 1.821(f) or 1.821(g) or 1.825(b) or 1.825(d).
For questions regarding compliance to these requirements, please contact:
For Rules Interpretation, call (703) 308-4216 For CRF Submission Help, call (703) 308-4212 Patentln Software Program Support
Technical Assistance703-287-0200 To Purchase PatentIn Software703-306-2600
PLEASE RETURN A COPY OF THIS NOTICE WITH YOUR REPLY